

**Phospho-PRAS40 (Thr246)
MAPmates™**

Cat. # 46-681

MILLIPLEX[®] MAP
Phospho PRAS40 (Thr246) MAPmates[™]
(100 Assay Points)

#46-681

INTRODUCTION

The MILLIPLEX MAP Phospho PRAS40 (Thr246) MAPmates pair is used in conjunction with the MILLIPLEX MAP Cell Signaling Buffer and Detection Kit (Catalog # 48-602) to detect the presence of phosphorylated PRAS40 (Thr246) in cell lysates using the Luminex[®] 100[™] IS, 200[™], or HTS system. Each MAPmates pair is ordered individually and may be combined for simultaneous multiplex analysis of cellular events. The MILLIPLEX MAP Cell Signaling Buffer and Detection Kit is ordered separately and consists of a common set of reagents needed for performing MAPmates assays. The detection assay is a rapid, convenient alternative to Western Blotting and immunoprecipitation procedures. Each kit contains sufficient reagents for one 96 well plate assay. The MILLIPLEX MAP HeLa Cell Lysate: Lambda Phosphatase treated included with this MAPmates pair may be utilized as a negative control for this target, while the HEK293 Cell Lysate: IGF Stimulated included with this MAPmates pair may be utilized as a positive control.

Important note: For a detailed protocol on Cell Signaling Detection Procedures please see the instruction booklet for the MILLIPLEX MAP Cell Signaling Buffer and Detection Kit (#48-602).

REAGENTS SUPPLIED	CATALOG NUMBER	LUMINEX BEAD #	VOLUME	QUANTITY
MILLIPLEX MAP Anti-PRAS40 Beads (20X)	42-681	55	131 μ L	1 tube
MILLIPLEX MAP Anti-phospho PRAS40 (Thr246), biotin (20X)	44-681	n/a	131 μ L	1 tube
MILLIPLEX MAP HeLa Cell Lysate: Lambda Phosphatase Treated	47-229	n/a	-----	1 vial
MILLIPLEX MAP HEK293 Cell Lysate: IGF Stimulated	47-228	n/a	-----	1 vial

STORAGE CONDITIONS UPON RECEIPT

- Recommended storage for kit components is 2 - 8°C.
- DO NOT FREEZE Streptavidin-Phycoerythrin.

This kit is for research purposes only.

Please read entire protocol before use.

It is important to use same assay incubation conditions throughout your study.

By purchasing this product, which contains fluorescently labeled microsphere beads authorized by Luminex Corporation ("Luminex"), you, the customer, acquire the right under Luminex's patent rights, if any, to use this product or any portion of this product, including without limitation the microsphere beads contained herein, only with Luminex's laser based fluorescent analytical test instrumentation marketed under the name of Luminex100, 200, HTS.

MATERIALS REQUIRED BUT NOT PROVIDED

Reagents

- MILLIPLEX MAP Cell Signaling Buffer and Detection Kit (Catalog # 48-602)
- Cell lysates or cell extracts harboring protein(s) of interest prepared using the MILLIPLEX MAP Cell Signaling Buffer and Detection Kit (Catalog # 48-602)
- Protease inhibitors (Millipore Catalog #20-201 or equivalent)
- Luminex Sheath Fluid (Luminex Catalog #40-50000)

Instrumentation / Materials

- Adjustable Pipettes with Tips capable of delivering 25 μ L to 1000 μ L
- Multichannel Pipettes capable of delivering 25 μ L to 200 μ L
- Reagent Reservoirs
- Polypropylene Microfuge Tubes
- Rubber Bands
- Absorbent Pads
- Laboratory Vortex Mixer
- Sonicator
- Titer Plate Shaker
- Vacuum Filtration Unit (Millipore Vacuum Manifold Catalog #MSVMHTS00 or equivalent with Millipore Vacuum Pump Catalog #WP6111560 or equivalent)
- Luminex 100™ IS, 200™, or HTS by Luminex Corporation
- Plate Stand (Millipore Catalog # MX-STAND)

SAFETY PRECAUTIONS

- All tissue components and biological materials should be handled as potentially hazardous. Follow universal precautions as established by the Centers for Disease Control and Prevention and by the Occupational Safety and Health Administration when handling and disposing of infectious agents.
- Sodium azide or Proclin has been added to some reagents as a preservative. Although the concentrations are low, sodium azide and Proclin may react with lead and copper plumbing to form highly explosive metal azides. On disposal, flush with a large volume of water to prevent azide build up.

TECHNICAL CONSIDERATIONS

- The suggested working range of protein concentration for the assay is 1 to 25 μ g of total protein/well (25 μ L/well at 40 to 1000 μ g/mL). A total protein amount of 10 μ g/ well is generally a good starting point for lysates for which target protein expression levels are unknown.
- The following MAPmates should not be multiplexed:
 - a. phospho-specific and total MAPmates pairs
 - b. pTyr and site-specific phospho MAPmates (the pTyr detect may generate false positives on the site-specific MAPmates)
 - c. Phospho MAPmates for a single target (Akt, STAT3, p53)

PREPARATION OF LYOPHILIZED MILLIPLEX^{MAP} CELL LYSATE

MILLIPLEX^{MAP} Cell Lysates as a positive and negative control

- Reconstitute each of the lyophilized cell lysates in 100 μ L of ultrapure water, for each vial this will yield 100 μ L of lysate at 2 mg/mL total protein.
- Gently vortex and incubate the reconstituted lysate for 5 min at RT (store on ice).
- Pipette **400** μ L MILLIPLEX^{MAP} Assay Buffer 1 to each of the reconstituted cell lysate vials. The cell lysate is now prepared for use in the MILLIPLEX^{MAP} Cell Signaling Assays.
- Combine prepared lysate (25 μ L/well) with 25 μ L Cell Signaling beads (step 4 of Immunoassay Protocol) and proceed with assay.

IMMUNOASSAY PROTOCOL

1. Dilute filtered lysates at least 1:4 (6.25 μL lysate + 18.75 μL Assay Buffer) in MILLIPLEX MAP Assay Buffer 1. The suggested working range of protein concentration for the assay is 1 to 25 μg of total protein/well (25 μL /well at 40 to 1,000 $\mu\text{g}/\text{ml}$).
2. Pre-wet filter plate with 25 μL /well of MILLIPLEX MAP Assay Buffer 1. Remove by vacuum filtration by placing the filter plate over a vacuum manifold and gently applying vacuum. Gently blot the bottom of the filter plate on a paper towel to remove excess liquid.
3. Gently mix the **1X** bead suspension with a pipette and sonicate for 10 seconds. Add 25 μL of 1X bead suspension to each well.
4. Add 25 μL of diluted cell lysate to each well and incubate overnight at 4°C (or 2 hours RT) on a plate shaker (600-800rpm) protected from light.
5. Remove the lysate by vacuum filtration.
6. Add 100 μL /well of MILLIPLEX MAP Assay Buffer 1. Remove buffer by vacuum filtration and gently blot the bottom of the filter plate on a paper towel.
7. Wash the beads a second time by repeating step 6.
8. Add 25 μL /well of **1X** MILLIPLEX MAP Biotinylated Reporter.
9. Incubate on a plate shaker for 1 hour at room temperature, protected from light.
10. Remove reporter by vacuum filtration and gently blot the bottom of the filter plate on a paper towel.
11. Add 25 μL of diluted (1:25) MILLIPLEX MAP Streptavidin-Phycoerythrin (SAPE).
12. Incubate on a plate shaker for 15 minutes at room temperature, protected from light.
13. **DO NOT REMOVE SAPE.** Add 25 μL of MILLIPLEX MAP Amplification Buffer to each well.

Add 25 μL Assay Buffer 1 per well



Remove buffer by vacuum.

- Add 25 μL 1X beads to wells
- Add 25 μL diluted cell lysate to appropriate wells



Incubate overnight at 4°C or 2 hour at RT with shaking; dark

Wash 2X with 100 μL Assay Buffer 1. Add 25 μL 1X reporter biotin.



Incubate 1 hr at RT with shaking; dark

Remove reporter and add 25 μL diluted SAPE.



Incubate 15 min at RT with shaking; dark

DO NOT REMOVE SAPE and add 25 μL Amplification buffer.

14. Incubate on a plate shaker for 15 minutes at room temperature, protected from light.
15. Remove MILLIPLEX MAP SAPE/Amplification buffer by vacuum filtration and gently blot the bottom of the filter plate on a paper towel.
16. Resuspend beads in 150 μ L of MILLIPLEX MAP Assay Buffer 1.
17. Analyze using the Luminex[®] system.



Incubate 15 min at RT with shaking; dark

Remove Streptavidin-PE/Amplification buffer and resuspend beads in 150 μ L Assay Buffer 1. Read results using appropriate Luminex[®] instrument.

EQUIPMENT SETTINGS

These specifications are for the Luminex₁₀₀ v.1.7 or Luminex100IS v2.1/2.2, Luminex₂₀₀ v2.3, xPONENT, and Luminex HTS. Luminex instruments with other software (e.g. MasterPlex, StarStation, LiquiChip, Bio-Plex, LABScan100) would need to follow instrument instructions for gate settings and additional specifications from the vendors.

Events	50 per bead
Sample Size	100 μ L
Gate Settings	8,000 to 15,000
Reporter Gain	Default (Low PMT)
Time Out	60 seconds
Bead Set	55

SINGLE AND MULTI-PLEX ANALYSIS

The recommended lysis and assay buffers for single or multi-plex analysis of Phospho PRAS40 (Thr246) MAPmates are MILLIPLEX MAP Lysis Buffer (Catalog # 43-040) and MILLIPLEX MAP Assay Buffer 1 (Catalog # 43-010). Both buffers are included in the MILLIPLEX MAP Cell Signaling Buffer and Detection Kit (Catalog # 48-602). For complete cell signaling assay and cell lysis protocols refer to the MILLIPLEX MAP Cell Signaling Buffer and Detection Kit instructions.

MAPmates which are listed in the MILLIPLEX MAP Cell Signaling Buffer and Detection Kit buffer selection chart as “not recommended” for the above buffer conditions must be assayed separately using appropriate buffer conditions.

Note: Phospho and Total MAPmates should not be multiplexed together.

REPRESENTATIVE DATA:

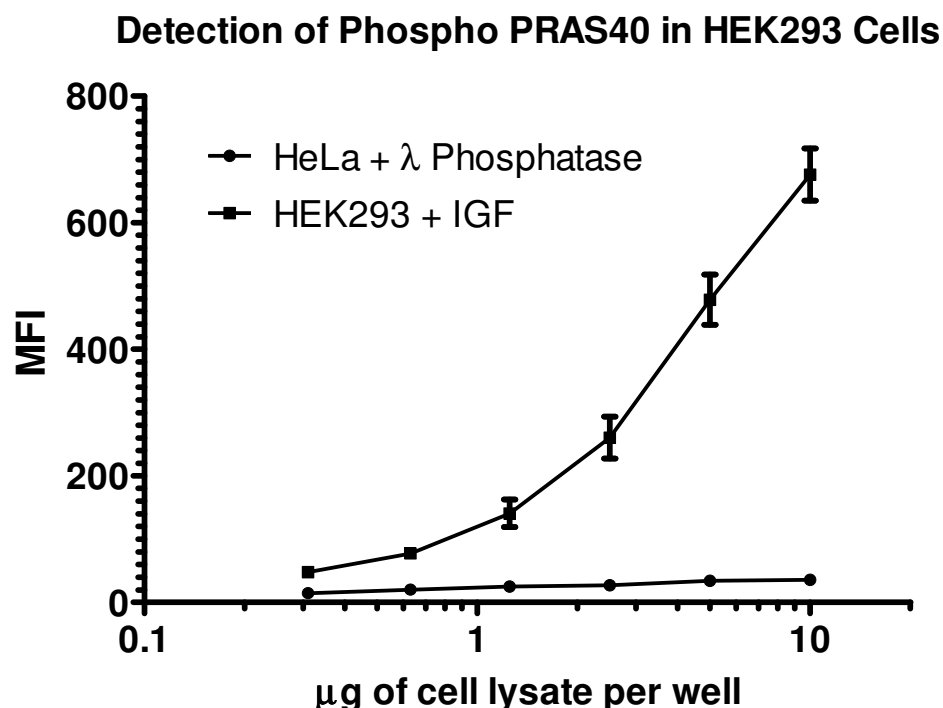


Figure 1. MILLIPLEX MAP detection of phosphorylated PRAS40 in HEK293 cell lysate. HEK293: IGF stimulated (#47-228) and HeLa: lambda phosphatase (#47-229) were prepared according to instructions. Increasing amounts of cell lysate were incubated overnight at 4°C with anti-PRAS40 capture beads. The beads were washed and probed with biotin labeled anti-phospho PRAS40, followed by incubation with streptavidin-PE and amplification buffer. The Median Fluorescent Intensity (MFI) in triplicate wells was measured using the Luminex® Instrument. This graph displays the MFI values obtained utilizing the Phospho PRAS40 MAPmates

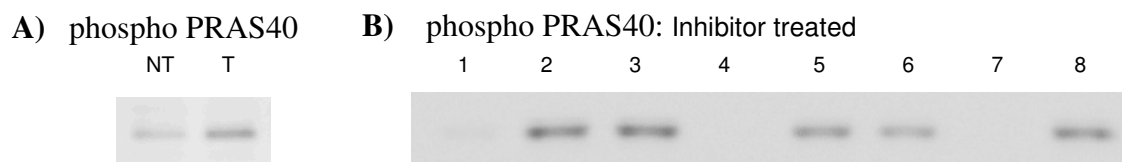


Figure 2. Immunoprecipitation/Western blot detection of phosphorylated PRAS40 in HEK293 cell lysate. **A)** HEK293 cells were grown to confluence and stimulated with or without 50ng/mL IGF for 15 minutes. 200 µg of HEK293 cell lysates (lysed in MILLIPLEX MAP Lysis Buffer with protease inhibitors) were mixed with capture antibody beads to immunoprecipitate phospho PRAS40 protein from non-treated (NT) or IGF1-treated (T) cell lysates. The immunoprecipitated proteins were separated on SDS-PAGE, transferred to nitrocellulose, and probed with biotin labeled phospho PRAS40 (Thr246) reporter antibody. The proteins were imaged using Streptavidin-HRP and chemiluminescence. **B)** HeLa (lane 1) and HEK293 cells (lanes 2 to 8) were grown to confluence and pretreated with 0.1µM wortmannin (lane 4), 10µM U0126 (lane 5), 0.1µM rapamycin (lane 6), 50µM LY294002 (lane 7) or 1µM Ro 31-8220 (lane 8) for 30 minutes before stimulating with 50ng/mL IGF (lanes 3 to 8) for 15 minutes. 20µg/well of cell lysates were separated by SDS-PAGE, transferred to nitrocellulose, and probed with biotin labeled phospho PRAS40 (Thr246) reporter antibody. Blots were incubated with Streptavidin-HRP and visualized via chemiluminescence.

ORDERING INFORMATION

To place an order:

FAX: (636) 441-8050

Include:

- Your name, telephone and/or fax number
- Customer account number
- Shipping and billing address
- Purchase order number
- Catalog number and description of product
- Quantity of kits

Toll-Free US: (800) MILLIPORE

Mail Orders: Millipore Corp.
6 Research Park Drive
St. Charles, Missouri 63304 U.S.A.

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Conditions of Sale

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